

CENTER FOR MEDICAL CANNABIS RESEARCH

2023 GENERAL SESSION

STATE OF UTAH

Chief Sponsor: Jennifer Dailey-Provost

Senate Sponsor: Evan J. Vickers

LONG TITLE

General Description:

This bill creates the Center for Medical Cannabis Research.

Highlighted Provisions:

This bill:

- ▶ defines terms;
- ▶ abolishes the Cannabis Research Review Board;
- ▶ creates the Center for Medical Cannabis Research (center) within the University of Utah; and
- ▶ establishes the center's duties.

Money Appropriated in this Bill:

None

Other Special Clauses:

None

Utah Code Sections Affected:

AMENDS:

- 4-41a-102**, as last amended by Laws of Utah 2022, Chapters 290, 452
- 26-61a-102**, as last amended by Laws of Utah 2022, Chapters 290, 452
- 26-61a-105**, as last amended by Laws of Utah 2022, Chapter 452
- 26-61a-201**, as last amended by Laws of Utah 2022, Chapters 198, 290 and 452
- 26-61a-703**, as last amended by Laws of Utah 2022, Chapter 97



28 ENACTS:

29 **53B-17-1401**, Utah Code Annotated 1953

30 **53B-17-1402**, Utah Code Annotated 1953

31 REPEALS:

32 **26-61-101**, as enacted by Laws of Utah 2017, Chapter 398

33 **26-61-102**, as last amended by Laws of Utah 2022, Chapter 452

34 **26-61-103**, as enacted by Laws of Utah 2017, Chapter 398

35 **26-61-201**, as last amended by Laws of Utah 2022, Chapter 452

36 **26-61-202**, as last amended by Laws of Utah 2022, Chapter 415



38 *Be it enacted by the Legislature of the state of Utah:*

39 Section 1. Section **4-41a-102** is amended to read:

40 **4-41a-102. Definitions.**

41 As used in this chapter:

42 (1) "Adulterant" means any poisonous or deleterious substance in a quantity that may
43 be injurious to health, including:

- 44 (a) pesticides;
- 45 (b) heavy metals;
- 46 (c) solvents;
- 47 (d) microbial life;
- 48 (e) toxins; or
- 49 (f) foreign matter.

50 ~~[(2) "Cannabis Research Review Board" means the Cannabis Research Review Board~~
51 ~~created in Section **26-61-201**.]~~

52 ~~[(3)]~~ (2) "Cannabis" means the same as that term is defined in Section **26-61a-102**.

53 ~~[(4)]~~ (3) "Cannabis concentrate" means:

- 54 (a) the product of any chemical or physical process applied to naturally occurring
55 biomass that concentrates or isolates the cannabinoids contained in the biomass; and
- 56 (b) any amount of a natural, derivative, or synthetic cannabinoid in the synthetic
57 cannabinoid's purified state.

58 ~~[(5)]~~ (4) "Cannabis cultivation byproduct" means any portion of a cannabis plant that is

59 not intended to be sold as a cannabis plant product.

60 ~~[(6)]~~ (5) "Cannabis cultivation facility" means a person that:

61 (a) possesses cannabis;

62 (b) grows or intends to grow cannabis; and

63 (c) sells or intends to sell cannabis to a cannabis cultivation facility, a cannabis
64 processing facility, or a medical cannabis research licensee.

65 ~~[(7)]~~ (6) "Cannabis cultivation facility agent" means an individual who:

66 (a) is an employee of a cannabis cultivation facility; and

67 (b) holds a valid cannabis production establishment agent registration card.

68 ~~[(8)]~~ (7) "Cannabis derivative product" means a product made using cannabis
69 concentrate.

70 ~~[(9)]~~ (8) "Cannabis plant product" means any portion of a cannabis plant intended to be
71 sold in a form that is recognizable as a portion of a cannabis plant.

72 ~~[(10)]~~ (9) "Cannabis processing facility" means a person that:

73 (a) acquires or intends to acquire cannabis from a cannabis production establishment;

74 (b) possesses cannabis with the intent to manufacture a cannabis product;

75 (c) manufactures or intends to manufacture a cannabis product from unprocessed
76 cannabis or a cannabis extract; and

77 (d) sells or intends to sell a cannabis product to a medical cannabis pharmacy or a
78 medical cannabis research licensee.

79 ~~[(11)]~~ (10) "Cannabis processing facility agent" means an individual who:

80 (a) is an employee of a cannabis processing facility; and

81 (b) holds a valid cannabis production establishment agent registration card.

82 ~~[(12)]~~ (11) "Cannabis product" means the same as that term is defined in Section
83 [26-61a-102](#).

84 ~~[(13)]~~ (12) "Cannabis production establishment" means a cannabis cultivation facility,
85 a cannabis processing facility, or an independent cannabis testing laboratory.

86 ~~[(14)]~~ (13) "Cannabis production establishment agent" means a cannabis cultivation
87 facility agent, a cannabis processing facility agent, or an independent cannabis testing
88 laboratory agent.

89 ~~[(15)]~~ (14) "Cannabis production establishment agent registration card" means a

90 registration card that the department issues that:

91 (a) authorizes an individual to act as a cannabis production establishment agent; and

92 (b) designates the type of cannabis production establishment for which an individual is
93 authorized to act as an agent.

94 [~~16~~] (15) "Community location" means a public or private elementary or secondary
95 school, a church, a public library, a public playground, or a public park.

96 [~~17~~] (16) "Cultivation space" means, quantified in square feet, the horizontal area in
97 which a cannabis cultivation facility cultivates cannabis, including each level of horizontal area
98 if the cannabis cultivation facility hangs, suspends, stacks, or otherwise positions plants above
99 other plants in multiple levels.

100 [~~18~~] (17) "Department" means the Department of Agriculture and Food.

101 [~~19~~] (18) "Derivative cannabinoid" means any cannabinoid that has been intentionally
102 created using a process to convert a naturally occurring cannabinoid into another cannabinoid.

103 [~~20~~] (19) "Family member" means a parent, step-parent, spouse, child, sibling,
104 step-sibling, uncle, aunt, nephew, niece, first cousin, mother-in-law, father-in-law,
105 brother-in-law, sister-in-law, son-in-law, daughter-in-law, grandparent, or grandchild.

106 [~~21~~] (20) (a) "Independent cannabis testing laboratory" means a person that:

107 (i) conducts a chemical or other analysis of cannabis or a cannabis product; or

108 (ii) acquires, possesses, and transports cannabis or a cannabis product with the intent to
109 conduct a chemical or other analysis of the cannabis or cannabis product.

110 (b) "Independent cannabis testing laboratory" includes a laboratory that the department
111 or a research university operates in accordance with Subsection 4-41a-201(14).

112 [~~22~~] (21) "Independent cannabis testing laboratory agent" means an individual who:

113 (a) is an employee of an independent cannabis testing laboratory; and

114 (b) holds a valid cannabis production establishment agent registration card.

115 [~~23~~] (22) "Industrial hemp waste" means:

116 (a) a cannabinoid concentrate; or

117 (b) industrial hemp biomass.

118 [~~24~~] (23) "Inventory control system" means a system described in Section 4-41a-103.

119 [~~25~~] (24) "Licensing board" or "board" means the Cannabis Production Establishment
120 Licensing Advisory Board created in Section 4-41a-201.1.

121 ~~[(26)]~~ (25) "Medical cannabis" means the same as that term is defined in Section
122 26-61a-102.

123 ~~[(27)]~~ (26) "Medical cannabis card" means the same as that term is defined in Section
124 26-61a-102.

125 ~~[(28)]~~ (27) "Medical cannabis pharmacy" means the same as that term is defined in
126 Section 26-61a-102.

127 ~~[(29)]~~ (28) "Medical cannabis pharmacy agent" means the same as that term is defined
128 in Section 26-61a-102.

129 ~~[(30)]~~ (29) "Medical cannabis research license" means a license that the department
130 issues to a research university for the purpose of obtaining and possessing medical cannabis for
131 academic research.

132 ~~[(31)]~~ (30) "Medical cannabis research licensee" means a research university that the
133 department licenses to obtain and possess medical cannabis for academic research, in
134 accordance with Section 4-41a-901.

135 ~~[(32)]~~ (31) "Medical cannabis treatment" means the same as that term is defined in
136 Section 26-61a-102.

137 ~~[(33)]~~ (32) "Medicinal dosage form" means the same as that term is defined in Section
138 26-61a-102.

139 ~~[(34)]~~ (33) "Qualified medical provider" means the same as that term is defined in
140 Section 26-61a-102.

141 ~~[(35)]~~ (34) "Qualified Production Enterprise Fund" means the fund created in Section
142 4-41a-104.

143 ~~[(36)]~~ (35) "Recommending medical provider" means the same as that term is defined
144 in Section 26-61a-102.

145 ~~[(37)]~~ (36) "Research university" means the same as that term is defined in Section
146 53B-7-702 and a private, nonprofit college or university in the state that:

- 147 (a) is accredited by the Northwest Commission on Colleges and Universities;
148 (b) grants doctoral degrees; and
149 (c) has a laboratory containing or a program researching a schedule I controlled
150 substance described in Section 58-37-4.

151 ~~[(38)]~~ (37) "State electronic verification system" means the system described in Section

152 26-61a-103.

153 [(39)] (38) "Synthetic cannabinoid" means any cannabinoid that:

154 (a) was chemically synthesized from starting materials other than a naturally occurring
155 cannabinoid; and

156 (b) is not a derivative cannabinoid.

157 [(40)] (39) "Tetrahydrocannabinol" or "THC" means the same as that term is defined in
158 Section 4-41-102.

159 [(41)] (40) "THC analog" means the same as that term is defined in Section 4-41-102.

160 [(42)] (41) "Total composite tetrahydrocannabinol" means all detectable forms of
161 tetrahydrocannabinol.

162 [(43)] (42) "Total tetrahydrocannabinol" or "total THC" means the same as that term is
163 defined in Section 4-41-102.

164 Section 2. Section 26-61a-102 is amended to read:

165 **26-61a-102. Definitions.**

166 As used in this chapter:

167 (1) "Active tetrahydrocannabinol" means THC, any THC analog, and
168 tetrahydrocannabinolic acid.

169 [(2) "~~Cannabis Research Review Board~~" means the ~~Cannabis Research Review Board~~
170 ~~created in Section 26-61-201.~~]

171 [(3)] (2) "Cannabis" means marijuana.

172 [(4)] (3) "Cannabis cultivation facility" means the same as that term is defined in
173 Section 4-41a-102.

174 [(5)] (4) "Cannabis processing facility" means the same as that term is defined in
175 Section 4-41a-102.

176 [(6)] (5) "Cannabis product" means a product that:

177 (a) is intended for human use; and

178 (b) contains cannabis or any tetrahydrocannabinol or THC analog in a total
179 concentration of 0.3% or greater on a dry weight basis.

180 [(7)] (6) "Cannabis production establishment" means the same as that term is defined
181 in Section 4-41a-102.

182 [(8)] (7) "Cannabis production establishment agent" means the same as that term is

183 defined in Section [4-41a-102](#).

184 ~~[(9)]~~ (8) "Cannabis production establishment agent registration card" means the same
185 as that term is defined in Section [4-41a-102](#).

186 ~~[(10)]~~ (9) "Community location" means a public or private elementary or secondary
187 school, a church, a public library, a public playground, or a public park.

188 ~~[(11)]~~ (10) "Conditional medical cannabis card" means an electronic medical cannabis
189 card that the department issues in accordance with Subsection [26-61a-201\(1\)\(b\)](#) to allow an
190 applicant for a medical cannabis card to access medical cannabis during the department's
191 review of the application.

192 ~~[(12)]~~ (11) "Controlled substance database" means the controlled substance database
193 created in Section [58-37f-201](#).

194 ~~[(13)]~~ (12) "Department" means the Department of Health and Human Services.

195 ~~[(14)]~~ (13) "Designated caregiver" means:

196 (a) an individual:

197 (i) whom an individual with a medical cannabis patient card or a medical cannabis
198 guardian card designates as the patient's caregiver; and

199 (ii) who registers with the department under Section [26-61a-202](#); or

200 (b) (i) a facility that an individual designates as a designated caregiver in accordance
201 with Subsection [26-61a-202\(1\)\(b\)](#); or

202 (ii) an assigned employee of the facility described in Subsection [26-61a-202\(1\)\(b\)\(ii\)](#).

203 ~~[(15)]~~ (14) "Directions of use" means recommended routes of administration for a
204 medical cannabis treatment and suggested usage guidelines.

205 ~~[(16)]~~ (15) "Dosing guidelines" means a quantity range and frequency of administration
206 for a recommended treatment of medical cannabis.

207 ~~[(17)]~~ (16) "Financial institution" means a bank, trust company, savings institution, or
208 credit union, chartered and supervised under state or federal law.

209 ~~[(18)]~~ (17) "Home delivery medical cannabis pharmacy" means a medical cannabis
210 pharmacy that the department authorizes, as part of the pharmacy's license, to deliver medical
211 cannabis shipments to a medical cannabis cardholder's home address to fulfill electronic orders
212 that the state central patient portal facilitates.

213 ~~[(19)]~~ (18) "Institutional review board" means an institutional review board that is

214 registered for human subject research by the United States Department of Health and Human
215 Services.

216 (19) "Inventory control system" means the system described in Section 4-41a-103.

217 (20) "Legal dosage limit" means an amount that:

218 (a) is sufficient to provide 30 days of treatment based on the dosing guidelines that the
219 relevant recommending medical provider or the state central patient portal or pharmacy
220 medical provider, in accordance with Subsection 26-61a-502(4) or (5), recommends; and

221 (b) may not exceed:

222 (i) for unprocessed cannabis in a medicinal dosage form, 113 grams by weight; and

223 (ii) for a cannabis product in a medicinal dosage form, a quantity that contains, in total,
224 greater than 20 grams of active tetrahydrocannabinol.

225 (21) "Legal use termination date" means a date on the label of a container of
226 unprocessed cannabis flower:

227 (a) that is 60 days after the date of purchase of the cannabis; and

228 (b) after which, the cannabis is no longer in a medicinal dosage form outside of the
229 primary residence of the relevant medical cannabis patient cardholder.

230 (22) "Limited medical provider" means an individual who:

231 (a) meets the recommending qualifications; and

232 (b) has no more than 15 patients with a valid medical cannabis patient card or
233 provisional patient card as a result of the individual's recommendation, in accordance with
234 Subsection 26-61a-106(1)(b).

235 (23) "Marijuana" means the same as that term is defined in Section 58-37-2.

236 (24) "Medical cannabis" means cannabis in a medicinal dosage form or a cannabis
237 product in a medicinal dosage form.

238 (25) "Medical cannabis card" means a medical cannabis patient card, a medical
239 cannabis guardian card, a medical cannabis caregiver card, or a conditional medical cannabis
240 card.

241 (26) "Medical cannabis cardholder" means:

242 (a) a holder of a medical cannabis card; or

243 (b) a facility or assigned employee, described in Subsection(14)(b), only:

244 (i) within the scope of the facility's or assigned employee's performance of the role of a

245 medical cannabis patient cardholder's caregiver designation under Subsection
246 26-61a-202(1)(b); and

247 (ii) while in possession of documentation that establishes:

248 (A) a caregiver designation described in Subsection 26-61a-202(1)(b);

249 (B) the identity of the individual presenting the documentation; and

250 (C) the relation of the individual presenting the documentation to the caregiver
251 designation.

252 (27) "Medical cannabis caregiver card" means an electronic document that a cardholder
253 may print or store on an electronic device or a physical card or document that:

254 (a) the department issues to an individual whom a medical cannabis patient cardholder
255 or a medical cannabis guardian cardholder designates as a designated caregiver; and

256 (b) is connected to the electronic verification system.

257 (28) "Medical cannabis courier" means a courier that:

258 (a) the department licenses in accordance with Section 26-61a-604; and

259 (b) contracts with a home delivery medical cannabis pharmacy to deliver medical
260 cannabis shipments to fulfill electronic orders that the state central patient portal facilitates.

261 (29) "Medical cannabis courier agent" means an individual who:

262 (a) is an employee of a medical cannabis courier; and

263 (b) who holds a valid medical cannabis courier agent registration card.

264 (30) (a) "Medical cannabis device" means a device that an individual uses to ingest or
265 inhale cannabis in a medicinal dosage form or a cannabis product in a medicinal dosage form.

266 (b) "Medical cannabis device" does not include a device that:

267 (i) facilitates cannabis combustion; or

268 (ii) an individual uses to ingest substances other than cannabis.

269 (31) "Medical cannabis guardian card" means an electronic document that a cardholder
270 may print or store on an electronic device or a physical card or document that:

271 (a) the department issues to the parent or legal guardian of a minor with a qualifying
272 condition; and

273 (b) is connected to the electronic verification system.

274 (32) "Medical cannabis patient card" means an electronic document that a cardholder
275 may print or store on an electronic device or a physical card or document that:

- 276 (a) the department issues to an individual with a qualifying condition; and
- 277 (b) is connected to the electronic verification system.
- 278 (33) "Medical cannabis pharmacy" means a person that:
- 279 (a) (i) acquires or intends to acquire medical cannabis or a cannabis product in a
- 280 medicinal dosage form from a cannabis processing facility or another medical cannabis
- 281 pharmacy or a medical cannabis device; or
- 282 (ii) possesses medical cannabis or a medical cannabis device; and
- 283 (b) sells or intends to sell medical cannabis or a medical cannabis device to a medical
- 284 cannabis cardholder.
- 285 (34) "Medical cannabis pharmacy agent" means an individual who:
- 286 (a) is an employee of a medical cannabis pharmacy; and
- 287 (b) who holds a valid medical cannabis pharmacy agent registration card.
- 288 (35) "Medical cannabis pharmacy agent registration card" means a registration card
- 289 issued by the department that authorizes an individual to act as a medical cannabis pharmacy
- 290 agent.
- 291 (36) "Medical cannabis shipment" means a shipment of medical cannabis or a medical
- 292 cannabis product that a home delivery medical cannabis pharmacy or a medical cannabis
- 293 courier delivers to a medical cannabis cardholder's home address to fulfill an electronic medical
- 294 cannabis order that the state central patient portal facilitates.
- 295 (37) "Medical cannabis treatment" means cannabis in a medicinal dosage form, a
- 296 cannabis product in a medicinal dosage form, or a medical cannabis device.
- 297 (38) (a) "Medicinal dosage form" means:
- 298 (i) for processed medical cannabis or a medical cannabis product, the following with a
- 299 specific and consistent cannabinoid content:
- 300 (A) a tablet;
- 301 (B) a capsule;
- 302 (C) a concentrated liquid or viscous oil;
- 303 (D) a liquid suspension that, after December 1, 2022, does not exceed 30 ml;
- 304 (E) a topical preparation;
- 305 (F) a transdermal preparation;
- 306 (G) a sublingual preparation;

307 (H) a gelatinous cube, gelatinous rectangular cuboid, or lozenge in a cube or
308 rectangular cuboid shape;

309 (I) a resin or wax; or

310 (J) an aerosol; or

311 (ii) for unprocessed cannabis flower, a container described in Section 4-41a-602 that:

312 (A) contains cannabis flowers in a quantity that varies by no more than 10% from the
313 stated weight at the time of packaging;

314 (B) at any time the medical cannabis cardholder transports or possesses the container in
315 public, is contained within an opaque bag or box that the medical cannabis pharmacy provides;
316 and

317 (C) is labeled with the container's content and weight, the date of purchase, the legal
318 use termination date, and after December 31, 2020, a barcode that provides information
319 connected to an inventory control system; and

320 (iii) a form measured in grams, milligrams, or milliliters.

321 (b) "Medicinal dosage form" includes a portion of unprocessed cannabis flower that:

322 (i) the medical cannabis cardholder has recently removed from the container described
323 in Subsection (38)(a)(ii) for use; and

324 (ii) does not exceed the quantity described in Subsection (38)(a)(ii).

325 (c) "Medicinal dosage form" does not include:

326 (i) any unprocessed cannabis flower outside of the container described in Subsection
327 (38)(a)(ii), except as provided in Subsection (38)(b);

328 (ii) any unprocessed cannabis flower in a container described in Subsection (38)(a)(ii)
329 after the legal use termination date;

330 (iii) a process of vaporizing and inhaling concentrated cannabis by placing the cannabis
331 on a nail or other metal object that is heated by a flame, including a blowtorch; or

332 (iv) a liquid suspension that is branded as a beverage.

333 (39) "Nonresident patient" means an individual who:

334 (a) is not a resident of Utah or has been a resident of Utah for less than 45 days;

335 (b) has a currently valid medical cannabis card or the equivalent of a medical cannabis
336 card under the laws of another state, district, territory, commonwealth, or insular possession of
337 the United States; and

338 (c) has been diagnosed with a qualifying condition as described in Section [26-61a-104](#).

339 (40) "Payment provider" means an entity that contracts with a cannabis production
340 establishment or medical cannabis pharmacy to facilitate transfers of funds between the
341 establishment or pharmacy and other businesses or individuals.

342 (41) "Pharmacy medical provider" means the medical provider required to be on site at
343 a medical cannabis pharmacy under Section [26-61a-403](#).

344 (42) "Provisional patient card" means a card that:

345 (a) the department issues to a minor with a qualifying condition for whom:

346 (i) a recommending medical provider has recommended a medical cannabis treatment;
347 and

348 (ii) the department issues a medical cannabis guardian card to the minor's parent or
349 legal guardian; and

350 (b) is connected to the electronic verification system.

351 (43) "Qualified medical provider" means an individual:

352 (a) who meets the recommending qualifications; and

353 (b) whom the department registers to recommend treatment with cannabis in a
354 medicinal dosage form under Section [26-61a-106](#).

355 (44) "Qualified Patient Enterprise Fund" means the enterprise fund created in Section
356 [26-61a-109](#).

357 (45) "Qualifying condition" means a condition described in Section [26-61a-104](#).

358 (46) "Recommend" or "recommendation" means, for a recommending medical
359 provider, the act of suggesting the use of medical cannabis treatment, which:

360 (a) certifies the patient's eligibility for a medical cannabis card; and

361 (b) may include, at the recommending medical provider's discretion, directions of use,
362 with or without dosing guidelines.

363 (47) "Recommending medical provider" means a qualified medical provider or a
364 limited medical provider.

365 (48) "Recommending qualifications" means that an individual:

366 (a) (i) has the authority to write a prescription;

367 (ii) is licensed to prescribe a controlled substance under Title 58, Chapter 37, Utah
368 Controlled Substances Act; and

369 (iii) possesses the authority, in accordance with the individual's scope of practice, to
370 prescribe a Schedule II controlled substance; and

371 (b) is licensed as:

372 (i) a podiatrist under Title 58, Chapter 5a, Podiatric Physician Licensing Act;

373 (ii) an advanced practice registered nurse under Title 58, Chapter 31b, Nurse Practice
374 Act;

375 (iii) a physician under Title 58, Chapter 67, Utah Medical Practice Act, or Title 58,
376 Chapter 68, Utah Osteopathic Medical Practice Act; or

377 (iv) a physician assistant under Title 58, Chapter 70a, Utah Physician Assistant Act.

378 (49) "State central patient portal" means the website the department creates, in
379 accordance with Section [26-61a-601](#), to facilitate patient safety, education, and an electronic
380 medical cannabis order.

381 (50) "State central patient portal medical provider" means a physician or pharmacist
382 that the department employs in relation to the state central patient portal to consult with
383 medical cannabis cardholders in accordance with Section [26-61a-602](#).

384 (51) "State electronic verification system" means the system described in Section
385 [26-61a-103](#).

386 (52) "Tetrahydrocannabinol" or "THC" means a substance derived from cannabis or a
387 synthetic equivalent as described in Subsection [58-37-4\(2\)\(a\)\(iii\)\(AA\)](#).

388 (53) "THC analog" means the same as that term is defined in Section [4-41-102](#).

389 (54) "Valid form of photo identification" means any of the following forms of
390 identification that is either current or has expired within the previous six months:

391 (a) a valid state-issued driver license or identification card;

392 (b) a valid United States federal-issued photo identification, including:

393 (i) a United States passport;

394 (ii) a United States passport card;

395 (iii) a United States military identification card; or

396 (iv) a permanent resident card or alien registration receipt card; or

397 (c) a passport that another country issued.

398 Section 3. Section [26-61a-105](#) is amended to read:

399 **[26-61a-105](#). Compassionate Use Board.**

400 (1) (a) The department shall establish a Compassionate Use Board consisting of:
401 (i) seven qualified medical providers that the executive director appoints and the
402 Senate confirms:
403 (A) who are knowledgeable about the medicinal use of cannabis;
404 (B) who are physicians licensed under Title 58, Chapter 67, Utah Medical Practice Act,
405 or Title 58, Chapter 68, Utah Osteopathic Medical Practice Act; and
406 (C) whom the appropriate board certifies in the specialty of neurology, pain medicine
407 and pain management, medical oncology, psychiatry, infectious disease, internal medicine,
408 pediatrics, or gastroenterology; and
409 (ii) as a nonvoting member and the chair of the Compassionate Use Board, the
410 executive director or the director's designee.
411 (b) In appointing the seven qualified medical providers described in Subsection (1)(a),
412 the executive director shall ensure that at least two have a board certification in pediatrics.
413 (2) (a) Of the members of the Compassionate Use Board that the executive director
414 first appoints:
415 (i) three shall serve an initial term of two years; and
416 (ii) the remaining members shall serve an initial term of four years.
417 (b) After an initial term described in Subsection (2)(a) expires:
418 (i) each term is four years; and
419 (ii) each board member is eligible for reappointment.
420 (c) A member of the Compassionate Use Board may serve until a successor is
421 appointed.
422 (3) Four members constitute a quorum of the Compassionate Use Board.
423 (4) A member of the Compassionate Use Board may receive:
424 (a) notwithstanding Section 63A-3-106, compensation or benefits for the member's
425 service; and
426 (b) travel expenses in accordance with Section 63A-3-107 and rules made by the
427 Division of Finance in accordance with Section 63A-3-107.
428 (5) The Compassionate Use Board shall:
429 (a) review and recommend for department approval a petition to the board regarding an
430 individual described in Subsection 26-61a-201(2)(a), a minor described in Subsection

431 26-61a-201(2)(c), or an individual who is not otherwise qualified to receive a medical cannabis
432 card to obtain a medical cannabis card for compassionate use, for the standard or a reduced
433 period of validity, if:

434 (i) for an individual who is not otherwise qualified to receive a medical cannabis card,
435 the individual's qualified medical provider is actively treating the individual for an intractable
436 condition that:

437 (A) substantially impairs the individual's quality of life; and

438 (B) has not, in the qualified medical provider's professional opinion, adequately
439 responded to conventional treatments;

440 (ii) the qualified medical provider:

441 (A) recommends that the individual or minor be allowed to use medical cannabis; and

442 (B) provides a letter, relevant treatment history, and notes or copies of progress notes
443 describing relevant treatment history including rationale for considering the use of medical
444 cannabis; and

445 (iii) the Compassionate Use Board determines that:

446 (A) the recommendation of the individual's qualified medical provider is justified; and

447 (B) based on available information, it may be in the best interests of the individual to
448 allow the use of medical cannabis;

449 (b) review and approve or deny the use of a medical cannabis device for an individual
450 described in Subsection 26-61a-201(2)(a)(i)(B) or a minor described in Subsection

451 26-61a-201(2)(c) if the individual's or minor's qualified medical provider recommends that the
452 individual or minor be allowed to use a medical cannabis device to vaporize the medical
453 cannabis treatment;

454 (c) unless no petitions are pending:

455 (i) meet to receive or review compassionate use petitions at least quarterly; and

456 (ii) if there are more petitions than the board can receive or review during the board's
457 regular schedule, as often as necessary;

458 (d) except as provided in Subsection (6), complete a review of each petition and
459 recommend to the department approval or denial of the applicant for qualification for a medical
460 cannabis card within 90 days after the day on which the board received the petition;

461 (e) consult with the department regarding the criteria described in Subsection (6); and

462 (f) report, before November 1 of each year, to the Health and Human Services Interim
463 Committee:

464 (i) the number of compassionate use recommendations the board issued during the past
465 year; and

466 (ii) the types of conditions for which the board recommended compassionate use.

467 (6) The department shall make rules, in consultation with the Compassionate Use
468 Board and in accordance with Title 63G, Chapter 3, Utah Administrative Rulemaking Act, to
469 establish a process and criteria for a petition to the board to automatically qualify for expedited
470 final review and approval or denial by the department in cases where, in the determination of
471 the department and the board:

472 (a) time is of the essence;

473 (b) engaging the full review process would be unreasonable in light of the petitioner's
474 physical condition; and

475 (c) sufficient factors are present regarding the petitioner's safety.

476 (7) (a) (i) The department shall review:

477 (A) any compassionate use for which the Compassionate Use Board recommends
478 approval under Subsection (5)(d) to determine whether the board properly exercised the board's
479 discretion under this section; and

480 (B) any expedited petitions the department receives under the process described in
481 Subsection (6).

482 (ii) If the department determines that the Compassionate Use Board properly exercised
483 the board's discretion in recommending approval under Subsection (5)(d) or that the expedited
484 petition merits approval based on the criteria established in accordance with Subsection (6), the
485 department shall:

486 (A) issue the relevant medical cannabis card; and

487 (B) provide for the renewal of the medical cannabis card in accordance with the
488 recommendation of the qualified medical provider described in Subsection (5)(a).

489 (b) (i) If the Compassionate Use Board recommends denial under Subsection (5)(d),
490 the individual seeking to obtain a medical cannabis card may petition the department to review
491 the board's decision.

492 (ii) If the department determines that the Compassionate Use Board's recommendation

493 for denial under Subsection (5)(d) was arbitrary or capricious:

494 (A) the department shall notify the Compassionate Use Board of the department's
495 determination; and

496 (B) the board shall reconsider the Compassionate Use Board's refusal to recommend
497 approval under this section.

498 (c) In reviewing the Compassionate Use Board's recommendation for approval or
499 denial under Subsection (5)(d) in accordance with this Subsection (7), the department shall
500 presume the board properly exercised the board's discretion unless the department determines
501 that the board's recommendation was arbitrary or capricious.

502 (8) Any individually identifiable health information contained in a petition that the
503 Compassionate Use Board or department receives under this section is a protected record in
504 accordance with Title 63G, Chapter 2, Government Records Access and Management Act.

505 ~~[(9) The Compassionate Use Board shall annually report the board's activity to the
506 Cannabis Research Review Board.]~~

507 Section 4. Section **26-61a-201** is amended to read:

508 **26-61a-201. Medical cannabis patient card -- Medical cannabis guardian card --**
509 **Conditional medical cannabis card -- Application -- Fees -- Studies.**

510 (1) (a) The department shall, within 15 days after the day on which an individual who
511 satisfies the eligibility criteria in this section or Section [26-61a-202](#) submits an application in
512 accordance with this section or Section [26-61a-202](#):

513 (i) issue a medical cannabis patient card to an individual described in Subsection

514 (2)(a);

515 (ii) issue a medical cannabis guardian card to an individual described in Subsection

516 (2)(b);

517 (iii) issue a provisional patient card to a minor described in Subsection (2)(c); and

518 (iv) issue a medical cannabis caregiver card to an individual described in Subsection

519 [26-61a-202](#)(4).

520 (b) (i) Beginning on the earlier of September 1, 2021, or the date on which the
521 electronic verification system is functionally capable of facilitating a conditional medical
522 cannabis card under this Subsection (1)(b), upon the entry of a recommending medical
523 provider's medical cannabis recommendation for a patient in the state electronic verification

524 system, either by the provider or the provider's employee or by a medical cannabis pharmacy
525 medical provider or medical cannabis pharmacy in accordance with Subsection
526 [26-61a-501](#)(10)(a), the department shall issue to the patient an electronic conditional medical
527 cannabis card, in accordance with this Subsection (1)(b).

528 (ii) A conditional medical cannabis card is valid for the lesser of:

529 (A) 60 days; or

530 (B) the day on which the department completes the department's review and issues a
531 medical cannabis card under Subsection (1)(a), denies the patient's medical cannabis card
532 application, or revokes the conditional medical cannabis card under Subsection (8).

533 (iii) The department may issue a conditional medical cannabis card to an individual
534 applying for a medical cannabis patient card for which approval of the Compassionate Use
535 Board is not required.

536 (iv) An individual described in Subsection (1)(b)(iii) has the rights, restrictions, and
537 obligations under law applicable to a holder of the medical cannabis card for which the
538 individual applies and for which the department issues the conditional medical cannabis card.

539 (2) (a) An individual is eligible for a medical cannabis patient card if:

540 (i) (A) the individual is at least 21 years old; or

541 (B) the individual is 18, 19, or 20 years old, the individual petitions the Compassionate
542 Use Board under Section [26-61a-105](#), and the Compassionate Use Board recommends
543 department approval of the petition;

544 (ii) the individual is a Utah resident;

545 (iii) the individual's recommending medical provider recommends treatment with
546 medical cannabis in accordance with Subsection (4);

547 (iv) the individual signs an acknowledgment stating that the individual received the
548 information described in Subsection (9); and

549 (v) the individual pays to the department a fee in an amount that, subject to Subsection
550 [26-61a-109](#)(5), the department sets in accordance with Section [63J-1-504](#).

551 (b) (i) An individual is eligible for a medical cannabis guardian card if the individual:

552 (A) is at least 18 years old;

553 (B) is a Utah resident;

554 (C) is the parent or legal guardian of a minor for whom the minor's qualified medical

555 provider recommends a medical cannabis treatment, the individual petitions the Compassionate
556 Use Board under Section 26-61a-105, and the Compassionate Use Board recommends
557 department approval of the petition;

558 (D) the individual signs an acknowledgment stating that the individual received the
559 information described in Subsection (9);

560 (E) pays to the department a fee in an amount that, subject to Subsection
561 26-61a-109(5), the department sets in accordance with Section 63J-1-504, plus the cost of the
562 criminal background check described in Section 26-61a-203; and

563 (F) the individual has not been convicted of a misdemeanor or felony drug distribution
564 offense under either state or federal law, unless the individual completed any imposed sentence
565 six months or more before the day on which the individual applies for a medical cannabis
566 guardian card.

567 (ii) The department shall notify the Department of Public Safety of each individual that
568 the department registers for a medical cannabis guardian card.

569 (c) (i) A minor is eligible for a provisional patient card if:

570 (A) the minor has a qualifying condition;

571 (B) the minor's qualified medical provider recommends a medical cannabis treatment
572 to address the minor's qualifying condition;

573 (C) one of the minor's parents or legal guardians petitions the Compassionate Use
574 Board under Section 26-61a-105, and the Compassionate Use Board recommends department
575 approval of the petition; and

576 (D) the minor's parent or legal guardian is eligible for a medical cannabis guardian card
577 under Subsection (2)(b) or designates a caregiver under Subsection (2)(d) who is eligible for a
578 medical cannabis caregiver card under Section 26-61a-202.

579 (ii) The department shall automatically issue a provisional patient card to the minor
580 described in Subsection (2)(c)(i) at the same time the department issues a medical cannabis
581 guardian card to the minor's parent or legal guardian.

582 (d) Beginning on the earlier of September 1, 2021, or the date on which the electronic
583 verification system is functionally capable of servicing the designation, if the parent or legal
584 guardian of a minor described in Subsections (2)(c)(i)(A) through (C) does not qualify for a
585 medical cannabis guardian card under Subsection (2)(b), the parent or legal guardian may

586 designate up to two caregivers in accordance with Subsection 26-61a-202(1)(c) to ensure that
587 the minor has adequate and safe access to the recommended medical cannabis treatment.

588 (3) (a) An individual who is eligible for a medical cannabis card described in
589 Subsection (2)(a) or (b) shall submit an application for a medical cannabis card to the
590 department:

591 (i) through an electronic application connected to the state electronic verification
592 system;

593 (ii) with the recommending medical provider; and

594 (iii) with information including:

595 (A) the applicant's name, gender, age, and address;

596 (B) the number of the applicant's valid form of photo identification;

597 (C) for a medical cannabis guardian card, the name, gender, and age of the minor
598 receiving a medical cannabis treatment under the cardholder's medical cannabis guardian card;
599 and

600 (D) for a provisional patient card, the name of the minor's parent or legal guardian who
601 holds the associated medical cannabis guardian card.

602 (b) The department shall ensure that a medical cannabis card the department issues
603 under this section contains the information described in Subsection (3)(a)(iii).

604 (c) (i) If a recommending medical provider determines that, because of age, illness, or
605 disability, a medical cannabis patient cardholder requires assistance in administering the
606 medical cannabis treatment that the recommending medical provider recommends, the
607 recommending medical provider may indicate the cardholder's need in the state electronic
608 verification system, either directly or, for a limited medical provider, through the order
609 described in Subsections 26-61a-106(1)(c) and (d).

610 (ii) If a recommending medical provider makes the indication described in Subsection
611 (3)(c)(i):

612 (A) the department shall add a label to the relevant medical cannabis patient card
613 indicating the cardholder's need for assistance;

614 (B) any adult who is 18 years old or older and who is physically present with the
615 cardholder at the time the cardholder needs to use the recommended medical cannabis
616 treatment may handle the medical cannabis treatment and any associated medical cannabis

617 device as needed to assist the cardholder in administering the recommended medical cannabis
618 treatment; and

619 (C) an individual of any age who is physically present with the cardholder in the event
620 of an emergency medical condition, as that term is defined in Section 31A-1-301, may handle
621 the medical cannabis treatment and any associated medical cannabis device as needed to assist
622 the cardholder in administering the recommended medical cannabis treatment.

623 (iii) A non-cardholding individual acting under Subsection (3)(c)(ii)(B) or (C) may not:

624 (A) ingest or inhale medical cannabis;

625 (B) possess, transport, or handle medical cannabis or a medical cannabis device outside
626 of the immediate area where the cardholder is present or with an intent other than to provide
627 assistance to the cardholder; or

628 (C) possess, transport, or handle medical cannabis or a medical cannabis device when
629 the cardholder is not in the process of being dosed with medical cannabis.

630 (4) To recommend a medical cannabis treatment to a patient or to renew a
631 recommendation, a recommending medical provider shall:

632 (a) before recommending or renewing a recommendation for medical cannabis in a
633 medicinal dosage form or a cannabis product in a medicinal dosage form:

634 (i) verify the patient's and, for a minor patient, the minor patient's parent or legal
635 guardian's valid form of identification described in Subsection (3)(a);

636 (ii) review any record related to the patient and, for a minor patient, the patient's parent
637 or legal guardian in:

638 (A) for a qualified medical provider, the state electronic verification system; and

639 (B) the controlled substance database created in Section 58-37f-201; and

640 (iii) consider the recommendation in light of the patient's qualifying condition, history
641 of substance use or opioid use disorder, and history of medical cannabis and controlled
642 substance use during an initial face-to-face visit with the patient; and

643 (b) state in the recommending medical provider's recommendation that the patient:

644 (i) suffers from a qualifying condition, including the type of qualifying condition; and

645 (ii) may benefit from treatment with cannabis in a medicinal dosage form or a cannabis
646 product in a medicinal dosage form.

647 (5) (a) Except as provided in Subsection (5)(b) or (c), a medical cannabis card that the

648 department issues under this section is valid for the lesser of:

649 (i) an amount of time that the recommending medical provider determines; or

650 (ii) (A) six months for the first issuance, and, except as provided in Subsection

651 (5)(a)(ii)(B), for a renewal; or

652 (B) for a renewal, one year if, after at least one year following the issuance of the

653 original medical cannabis card, the recommending medical provider determines that the patient

654 has been stabilized on the medical cannabis treatment and a one-year renewal period is

655 justified.

656 (b) (i) A medical cannabis card that the department issues in relation to a terminal

657 illness described in Section 26-61a-104 expires after one year.

658 (ii) The recommending medical provider may revoke a recommendation that the

659 provider made in relation to a terminal illness described in Section 26-61a-104 if the medical

660 cannabis cardholder no longer has the terminal illness.

661 (c) A medical cannabis card that the department issues in relation to acute pain as

662 described in Section 26-61a-104 expires 30 days after the day on which the department first

663 issues a conditional or full medical cannabis card.

664 (6) (a) A medical cannabis patient card or a medical cannabis guardian card is

665 renewable if:

666 (i) at the time of renewal, the cardholder meets the requirements of Subsection (2)(a) or

667 (b); or

668 (ii) the cardholder received the medical cannabis card through the recommendation of

669 the Compassionate Use Board under Section 26-61a-105.

670 (b) The recommending medical provider who made the underlying recommendation

671 for the card of a cardholder described in Subsection (6)(a) may renew the cardholder's card

672 through phone or video conference with the cardholder, at the recommending medical

673 provider's discretion.

674 (c) Before having access to a renewed card, a cardholder under Subsection (2)(a) or (b)

675 shall pay to the department a renewal fee in an amount that:

676 (i) subject to Subsection 26-61a-109(5), the department sets in accordance with Section

677 63J-1-504; and

678 (ii) may not exceed the cost of the relatively lower administrative burden of renewal in

679 comparison to the original application process.

680 (d) If a minor meets the requirements of Subsection (2)(c), the minor's provisional
681 patient card renews automatically at the time the minor's parent or legal guardian renews the
682 parent or legal guardian's associated medical cannabis guardian card.

683 (7) (a) A cardholder under this section shall carry the cardholder's valid medical
684 cannabis card with the patient's name.

685 (b) (i) A medical cannabis patient cardholder or a provisional patient cardholder may
686 purchase, in accordance with this chapter and the recommendation underlying the card,
687 cannabis in a medicinal dosage form, a cannabis product in a medicinal dosage form, or a
688 medical cannabis device.

689 (ii) A cardholder under this section may possess or transport, in accordance with this
690 chapter and the recommendation underlying the card, cannabis in a medicinal dosage form, a
691 cannabis product in a medicinal dosage form, or a medical cannabis device.

692 (iii) To address the qualifying condition underlying the medical cannabis treatment
693 recommendation:

694 (A) a medical cannabis patient cardholder or a provisional patient cardholder may use
695 cannabis in a medicinal dosage form, a medical cannabis product in a medicinal dosage form,
696 or a medical cannabis device; and

697 (B) a medical cannabis guardian cardholder may assist the associated provisional
698 patient cardholder with the use of cannabis in a medicinal dosage form, a medical cannabis
699 product in a medicinal dosage form, or a medical cannabis device.

700 (8) The department may revoke a medical cannabis card that the department issues
701 under this section if the cardholder:

702 (a) violates this chapter; or

703 (b) is convicted under state or federal law of, after March 17, 2021, a drug distribution
704 offense.

705 (9) The department shall establish by rule, in accordance with Title 63G, Chapter 3,
706 Utah Administrative Rulemaking Act, a process to provide information regarding the following
707 to an individual receiving a medical cannabis card:

708 (a) risks associated with medical cannabis treatment;

709 (b) the fact that a condition's listing as a qualifying condition does not suggest that

710 medical cannabis treatment is an effective treatment or cure for that condition, as described in
711 Subsection 26-61a-104(1); and

712 (c) other relevant warnings and safety information that the department determines.

713 (10) The department may establish procedures by rule, in accordance with Title 63G,
714 Chapter 3, Utah Administrative Rulemaking Act, to implement the application and issuance
715 provisions of this section.

716 (11) (a) On or before September 1, 2021, the department shall establish by rule, in
717 accordance with Title 63G, Chapter 3, Utah Administrative Rulemaking Act, a process to allow
718 an individual from another state to register with the department in order to purchase medical
719 cannabis or a medical cannabis device from a medical cannabis pharmacy while the individual
720 is visiting the state.

721 (b) The department may only provide the registration process described in Subsection
722 (11)(a):

723 (i) to a nonresident patient; and

724 (ii) for no more than two visitation periods per calendar year of up to 21 calendar days
725 per visitation period.

726 (12) (a) A person may submit to the department a request to conduct a research study
727 using medical cannabis cardholder data that the state electronic verification system contains.

728 (b) The department shall review a request described in Subsection (12)(a) to determine
729 whether an institutional review board~~[, as that term is defined in Section 26-61-102,]~~ could
730 approve the research study.

731 (c) At the time an individual applies for a medical cannabis card, the department shall
732 notify the individual:

733 (i) of how the individual's information will be used as a cardholder;

734 (ii) that by applying for a medical cannabis card, unless the individual withdraws
735 consent under Subsection (12)(d), the individual consents to the use of the individual's
736 information for external research; and

737 (iii) that the individual may withdraw consent for the use of the individual's
738 information for external research at any time, including at the time of application.

739 (d) An applicant may, through the medical cannabis card application, and a medical
740 cannabis cardholder may, through the state central patient portal, withdraw the applicant's or

741 cardholder's consent to participate in external research at any time.

742 (e) The department may release, for the purposes of a study described in this
743 Subsection (12), information about a cardholder under this section who consents to participate
744 under Subsection (12)(c).

745 (f) If an individual withdraws consent under Subsection (12)(d), the withdrawal of
746 consent:

747 (i) applies to external research that is initiated after the withdrawal of consent; and

748 (ii) does not apply to research that was initiated before the withdrawal of consent.

749 (g) The department may establish standards for a medical research study's validity, by
750 rule made in accordance with Title 63G, Chapter 3, Utah Administrative Rulemaking Act.

751 (13) The department shall record the issuance or revocation of a medical cannabis card
752 under this section in the controlled substance database.

753 Section 5. Section **26-61a-703** is amended to read:

754 **26-61a-703. Report.**

755 (1) By the November interim meeting each year beginning in 2020, the department
756 shall report to the Health and Human Services Interim Committee on:

757 (a) the number of applications and renewal applications filed for medical cannabis
758 cards;

759 (b) the number of qualifying patients and designated caregivers;

760 (c) the nature of the debilitating medical conditions of the qualifying patients;

761 (d) the age and county of residence of cardholders;

762 (e) the number of medical cannabis cards revoked;

763 (f) the number of practitioners providing recommendations for qualifying patients;

764 (g) the number of license applications and renewal license applications received;

765 (h) the number of licenses the department has issued in each county;

766 (i) the number of licenses the department has revoked;

767 (j) the quantity of medical cannabis shipments that the state central patient portal
768 facilitates;

769 (k) the number of overall purchases of medical cannabis and medical cannabis products
770 from each medical cannabis pharmacy;

771 (l) the expenses incurred and revenues generated from the medical cannabis program;

772 and

773 (m) an analysis of product availability in medical cannabis pharmacies.

774 (2) The report shall include information provided by the Center for Medical Cannabis
775 Research described in Section 53B-17-1402.

776 [~~2~~] (3) The department may not include personally identifying information in the
777 report described in this section.

778 [~~3~~] (4) During the 2022 legislative interim, the department shall report to the working
779 group described in Section 36-12-8.2 as requested by the working group.

780 Section 6. Section 53B-17-1401 is enacted to read:

781 **CHAPTER 17. UNIVERSITY OF UTAH**

782 **Part 14. Center for Medical Cannabis Research**

783 **53B-17-1401. Definitions.**

784 As used in this part:

785 (1) "Academic research cannabis license" means the license described in Title 4,
786 Chapter 41a, Part 9, Academic Medical Cannabis Research.

787 (2) "Cannabis" means the same as that term is defined in Section 26-61a-102.

788 (3) "Cannabis cultivation facility" means the same as that term is defined in Section
789 4-41a-102.

790 (4) "Cannabis product" means the same as that term is defined in Section 26-61a-102.

791 (5) "Center" means the Center for the Medical Cannabis Research created in Section
792 53B-17-1402.

793 (6) "Eligible institution" means an institution of higher education that:

794 (a) is located in Utah; and

795 (b) has or will obtain an academic research cannabis license.

796 (7) "Medical cannabis patient card" means the same as that term is defined in Section
797 26-61a-102.

798 Section 7. Section 53B-17-1402 is enacted to read:

799 **53B-17-1402. Center creation -- Duties.**

800 (1) There is created the Center for Medical Cannabis Research within the University of
801 Utah.

802 (2) The center:

- 803 (a) shall seek state, federal, and private funds to award grants for medical cannabis
804 research;
- 805 (b) shall facilitate and support funding for research related to the health effects,
806 including the potential risks or side effects, of the use of cannabis products;
- 807 (c) shall facilitate and support funding for research related to the efficacy and potential
808 health effects of various cannabis delivery methods, including vaporizing, ingesting, topical
809 application, and combustion;
- 810 (d) shall support researchers in applying for and securing federal and private research
811 grant funding for expanding medical cannabis research;
- 812 (e) shall review current and future cannabis research literature, clinical studies, and
813 clinical trials;
- 814 (f) shall educate medical providers, lawmakers, and the public about medical cannabis
815 research advances;
- 816 (g) shall, if requested, consult with researchers and eligible institutions seeking to
817 conduct medical cannabis research regarding legal implications of the research under state and
818 federal law;
- 819 (h) shall monitor, to the extent that appropriate and sufficient data are available, patient
820 outcomes in any state with a medicinal cannabis program;
- 821 (i) may coordinate, share knowledge, and share best practices with a state:
822 (i) that has a medical cannabis program; and
823 (ii) is conducting cannabis research;
- 824 (j) may award or facilitate funding for grants to an eligible institution for medical
825 cannabis research, including research regarding the growing of a medical-grade cannabis plant
826 that is used for a cannabis product;
- 827 (k) shall support a licensed cannabis cultivation facility to provide medical-grade
828 cannabis products for research;
- 829 (l) shall make any research conducted by the center publicly available;
- 830 (m) shall maintain a catalog of all published scientific reports based on projects funded
831 or managed by the center;
- 832 (n) shall ensure that an individual who agrees to use a cannabis product as part of a
833 research project conducted by the center or a grantee has:

- 834 (i) a valid medical cannabis patient card from the state; or
- 835 (ii) if included in the research project as a resident of another state, the equivalent of a
- 836 medical cannabis patient card under the laws of another state, district, territory,
- 837 commonwealth, or insular possession of the United States;
- 838 (o) shall obtain an academic research cannabis license;
- 839 (p) may apply for, or assist an eligible institution to apply for, a federal cannabis
- 840 cultivation registration to locate a cannabis cultivation site in Utah; and
- 841 (q) for the report described in Section 26-61a-703, shall provide information to the
- 842 Department of Health and Human Services describing:
- 843 (i) all research projects that are funded by a grant awarded by the center, including
- 844 which institution received the grant; and
- 845 (ii) all research projects conducted by the center.
- 846 (3) The University of Utah shall provide staff for the center.

847 Section 8. **Repealer.**

848 This bill repeals:

849 Section **26-61-101, Title.**

850 Section **26-61-102, Definitions.**

851 Section **26-61-103, Institutional review board -- Approved study of cannabis, a**

852 **cannabinoid product, or an expanded cannabinoid product.**

853 Section **26-61-201, Cannabis Research Review Board.**

854 Section **26-61-202, Duties.**